

AMENDMENTS TO THE SPECIFICATION**IN THE SPECIFICATION:****Page 18**

Please amend the Specification on page 18 beginning at line 15 as follows:

An aqueous solution of 100 µg/mL GFP expression plasmid (100 µL) was added to each well of a 24-well cell culture plate, and dried by blowing cool air. Human ~~adrenogenie~~ embryonic kidney epithelial cell line, 293 cells (ATCC: Cell Biology Collection) were suspended into the DMEM medium (Sigma) containing 10% fetal bovine serum (FBS), and seeded at 2.5×10^4 cells (500 µL)/well. Immediately after the cells were seeded, an aqueous solution of 1.7 M calcium chloride (0, 1.5, 2.5, 3.5, 5.0, 6.5, or 8.0 µL) was added and the plate was stirred to mix uniformly. The final concentration of calcium chloride in each medium was 1.8 mM, 7.1 mM, 10.2 mM, 14.2 mM, 19.5 mM, 24.8 mM, or 30.1 mM. On day 2 after seeding, cells were observed with fluorescence microscope, and the cells expressing GFP were counted to calculate the transfer efficiency.

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Please amend the Specification on page 21 beginning at line 30 and continuing onto page 22 as follows:

A solution containing complexes was prepared by mixing equal volumes of an aqueous solution of GFP expression plasmid (200 µg/mL) and an aqueous solution of 0.016 % atelocollagen. The complex solution (100 µL) was added to each well of a 24-well cell culture

plate, and dried by blowing cool air. Human adrenogenie embryonic kidney epithelial cell line, 293 cells or human cervical cancer-derived epithelial cell line, HeLa cells (ATCC: Cell Biology Collection) were suspended into the DMEM medium (Sigma) containing 10% fatal bovine serum (FBS), and seeded at 2.5×10^4 cells (500 μ L)/well. Immediately after the cells were seeded, an aqueous solution of 1.7 M calcium chloride (5.0 μ L) was added and the plate was stirred to mix uniformly. On day 2 after seeding, cells were observed with fluorescence microscope, and the cells expressing GFP were counted to calculate the transfer efficiency.